

Structure attributes must be viewed using STN Express query preparation.

=> s 11 full  
L3 27 SEA SSS FUL L1

=> file ca

=> s 13  
L4 1 L3

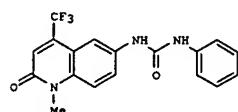
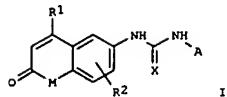
=> d ibib abs fhitstr

L4 ANSWER 1 OF 1 CA COPYRIGHT 2005 ACS on STN  
 ACCESSION NUMBER: 141:207071 CA  
 TITLE: Preparation of quinoline and chromene urea and thiourea derivatives as androgen receptor antagonists  
 INVENTOR(S): Du, Daniel Yunlong; Procter, Martin James; Fyfe, Matthew Colin Thor; Shah, Vilasben Kanji; Williams, Geoffrey Martyn; Schofield, Karen Lesley  
 PATENT ASSIGNEE(S): Warner-Lambert Company LLC, USA  
 SOURCE: PCT Int. Appl. 37 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004072044	A2	20040826	WO 2004-IB295.	20040130
WO 2004072044	A3	20041111		
W: AE, AE, AG, AL, AL, AM, AM, AM, AT, AT, AU, A2, A2, BA, BB, BG, BG, BR, BR, BW, BY, BY, BZ, BZ, CA, CH, CN, CN, CO, CO, CR, CR, CU, CU, CZ, CZ, DE, DE, DK, DK, DM, DZ, EC, EC, EE, EE, EG, ES, ES, FI, FI, GB, GD, GE, GE, GH, GM, HR, HR, HU, HU, ID, IL, IN, IS, JP, JP, KE, KE, KG, KG, KP, KP, KR, KR, KZ, KZ, KZ, LC, LK, LR, LS, LS, LT, LU, LV, MA, MD, MD, MG, MK, MN, MW, MX, MX, MZ, MZ, NA, NI, RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GO, GW, ML, MR, NE, SN, TD, TG, BF, BJ, CF, CG, CI, CM, GA, GN, GO, GW, ML, MR, NE, SN, TD, TG				
US 2004266820	A1	20041230	US 2004-775464	20040210
PRIORITY APPN. INFO.:			US 2003-446409P	P 20030211

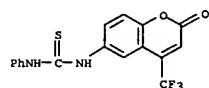
OTHER SOURCE(S): MARPAT 141:207071  
 GI

L4 ANSWER 1 OF 1 CA COPYRIGHT 2005 ACS on STN (Continued)



AB The title compds. I [M = N2 or O; Z = H, alkyl; R1 = H, alkyl, optionally substituted with one or more halogens, or alkoxy, optionally substituted with one or more halogens; R2 = absent or may represent up to 2 substituents selected from halo, CN, OH, alkyl, alkenyl, alkynyl, alkoxy, etc.; X = O or S; A = H, alkyl, alkenyl, alkynyl, etc.] were prepared as androgen receptor antagonists for the treatment of alopecia, acne, oily skin, prostate cancer, hirsutism, and benign prostate hyperplasia. For example, reaction of 6-amino-1-methyl-4-trifluoromethyl-1H-quinoline-2-one (preparation given) with Ph isocyanate yielded compound II.

IT 743467-64-3P  
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)  
 (Preparation of urea and thiourea derivs. as androgen receptor antagonists)  
 RN 743467-64-3 CA  
 CN Thiourea, N-[2-oxo-4-(trifluoromethyl)-2H-1-benzopyran-6-yl]-N'-phenyl- (9CI) (CA INDEX NAME)



10/775464

=> file marpat

=> s 11 full

FULL SEARCH INITIATED 14:44:13 FILE 'MARPAT'  
FULL SCREEN SEARCH COMPLETED - 951 TO ITERATE

100.0% PROCESSED 951 ITERATIONS  
SEARCH TIME: 00.00.02

1 ANSWERS

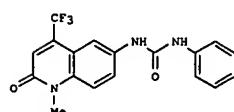
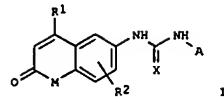
L5 1 SEA SSS FUL L1

L5 ANSWER 1 OF 1 MARPAT COPYRIGHT 2005 ACS on STN  
 ACCESSION NUMBER: 141:207071 MARPAT  
 TITLE: Preparation of quinoline and chromene ureas and thiourea derivatives as androgen receptor antagonists  
 INVENTOR(S): Du, Daniel Yunlong; Procter, Martin James; Fyfe, Matthew Colin Thor; Shah, Vilasaben Kanji; Williams, Geoffrey Martyn; Schofield, Karen Lesley  
 PATENT ASSIGNEE(S): Warner-Lambert Company LLC, USA  
 SOURCE: PCT Int. Appl., 37 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004072044	A2	20040826	WO 2004-IB295	20040130
WO 2004072044	A3	20041111		
W:	AE, AE, AG, AL, AL, AM, AM, AM, AT, AT, AU, AZ, A2, BA, BB, BG, BG, BR, BR, BW, BY, BZ, BZ, CA, CH, CN, CO, CO, CR, CR, CU, CU, CZ, CZ, DE, DE, DK, DK, DM, DZ, EC, EC, EE, EE, EG, ES, ES, FI, FI, GB, GD, GE, GE, GH, GM, HR, HR, HU, HU, ID, IL, IN, IS, JP, JP, KE, KE, KG, KG, KP, KP, KR, KR, KZ, KZ, LC, LK, LR, LS, LS, LT, LU, LV, MA, MD, MD, MG, MK, MN, MW, MX, MX, MZ, MZ, NA, NI, RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BF, BJ, CF, CG, CI, CM, GA, GN, US 2004266820 A1 20041230 US 2004-775464 20040210 US 2003-446409P 20030211			

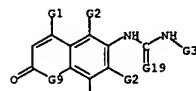
PRIORITY APPLN. INFO.: GI

L5 ANSWER 1 OF 1 MARPAT COPYRIGHT 2005 ACS on STN (Continued)



AB: The title compds. I [M = NZ or O; Z = H, alkyl; R1 = H, alkyl, optionally substituted with one or more halogens, or alkoxy, optionally substituted with one or more halogens; R2 = absent or may represent up to 2 substituents selected from halo, CN, OH, alkyl, Alkenyl, alkynyl, alkoxy, etc.]; X = O or S; A = H, alkyl, alkenyl, alkynyl, etc. were prepared as androgen receptor antagonists for the treatment of alopecia, acne, oily skin, prostate cancer, hirsutism, and benign prostate hyperplasia. For example, reaction of 6-amino-1-methyl-4-trifluoromethyl-1H-quinoline-2-one (preparation given) with Ph isocyanate yielded compound II.

MSTR 1



G1 = CF3  
 G9 = O  
 G19 = O

Patent location: claim 1

10/775464

=> d his

(FILE 'HOME' ENTERED AT 14:40:36 ON 04 OCT 2005)

FILE 'REGISTRY' ENTERED AT 14:43:24 ON 04 OCT 2005

L1 STRUCTURE uploaded

L2 1 S L1 SAM

L3 27 S L1 FULL

FILE 'CA' ENTERED AT 14:43:51 ON 04 OCT 2005

L4 1 S L3

FILE 'MARPAT' ENTERED AT 14:44:09 ON 04 OCT 2005

L5 1 S L1 FULL

=>

---Logging off of STN---

=>

Executing the logoff script...

=> LOG Y

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
----------------------	------------	-------

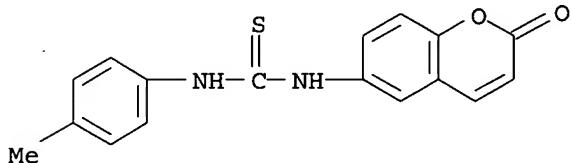
STN INTERNATIONAL LOGOFF AT 14:44:45 ON 04 OCT 2005

'/Q'. The saved name for an answer set must end with '/A'. The saved name for an L# list must end with '/L'. SDI request names must end with '/S'. To see a list of all saved query, answer set,, and L# list names for this loginid, enter "DISPLAY SAVED" at an arrow prompt (=>). Enter "DISPLAY SAVED/S" to see a list of SDI request names. Enter "DISPLAY SAVED/B" to see a list of BATCH search requests.

=> s 101444-67-1/rn  
L3 1 101444-67-1/RN

=> d

L3 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2005 ACS on STN  
RN 101444-67-1 REGISTRY  
ED Entered STN: 12 Apr 1986  
CN Coumarin, 6-(2-thio-3-p-tolylureido)- (6CI) (CA INDEX NAME)  
FS 3D CONCORD  
MF C17 H14 N2 O2 S  
SR CAOLD  
LC STN Files: BEILSTEIN\*, CA, CAOLD, CAPLUS  
(\*File contains numerically searchable property data)

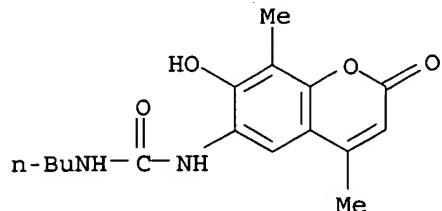


\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE)  
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)  
1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

=>

ANSWER 1 OF 1 REGISTRY COPYRIGHT 2005 ACS on STN  
 RN 176913-89-6 REGISTRY  
 ED Entered STN: 04 Jun 1996  
 CN Urea, N-butyl-N'-(7-hydroxy-4,8-dimethyl-2-oxo-2H-1-benzopyran-6-yl)-  
 (9CI) (CA INDEX NAME)  
 FS 3D CONCORD  
 MF C16 H20 N2 O4  
 SR CA  
 LC STN Files: CA, CAPLUS, TOXCENTER



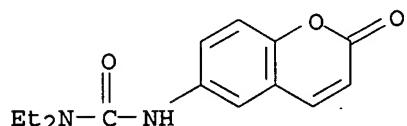
\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE)  
 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> s 6513-61-7/rn  
 L2 1 6513-61-7/RN

=> d

L2 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2005 ACS on STN  
 RN 6513-61-7 REGISTRY  
 ED Entered STN: 16 Nov 1984  
 CN Coumarin, 6-(3,3-diethylureido)- (7CI, 8CI) (CA INDEX NAME)  
 FS 3D CONCORD  
 MF C14 H16 N2 O3  
 LC STN Files: BEILSTEIN\*, CA, CAOLD, CAPLUS  
 (\*File contains numerically searchable property data)



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE)  
 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)  
 1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

=> d 101444-67-1/rn  
 '101444-67-1' MUST END IN '/Q', '/A', '/L', '/S' OR '/B'  
 The saved name for a query (or structure or screen set) must end with